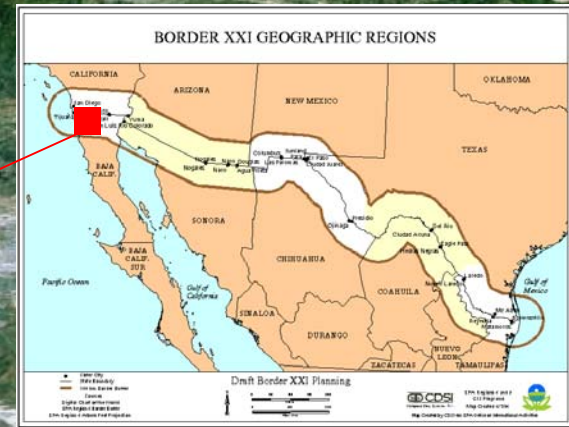


Research Translation

GOAL: “Move research findings/technology into decision-making processes through communication” (Dr. William Suk)

WHY?to strengthen public, private, tribal and/or non-profit efforts aimed at improving human health and the environmental systems upon which human health depends.

HOW?create innovative knowledge-action collaboratives that can holistically integrate biomedical, geochemical, engineering, social and information sciences in applied contexts.



San Diego-Tijuana City-Region

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Community Outreach Core

Dr. Keith Pezzoli, Core Leader; Dr. Richard Marciano, Co-Leader; Dr. David Pellow, Co-Leader
([Core Leader and Co-Leader bios](#))

Integrating Superfund Science and Traditional Environmental Knowledge: A Tribal Regional Workbench Approach



Tribal Partners

Michael Connolly, Laguna Resources, Campo Indian Reservation
Phil Green, Environmental Coordinator, Campo Tribal EPA
Ralph Goff, Tribal Chairman, Campo Indian Reservation
David Conrad, Executive Director, National Tribal Environmental Council (NTEC)
Lisa Gover, NTEC and Superfund Working Group

Dean Mike, Tribal Chairman, 29 Palms Tribal EPA

Dr. Marshall Cheung, Environmental Coordinator, 29 Palms Tribal EPA

Ken Bailey, Bureau of Indian Affairs



**Superfund sites/ w
50 mile buffer**



**US EPA Region 9
Indian lands**



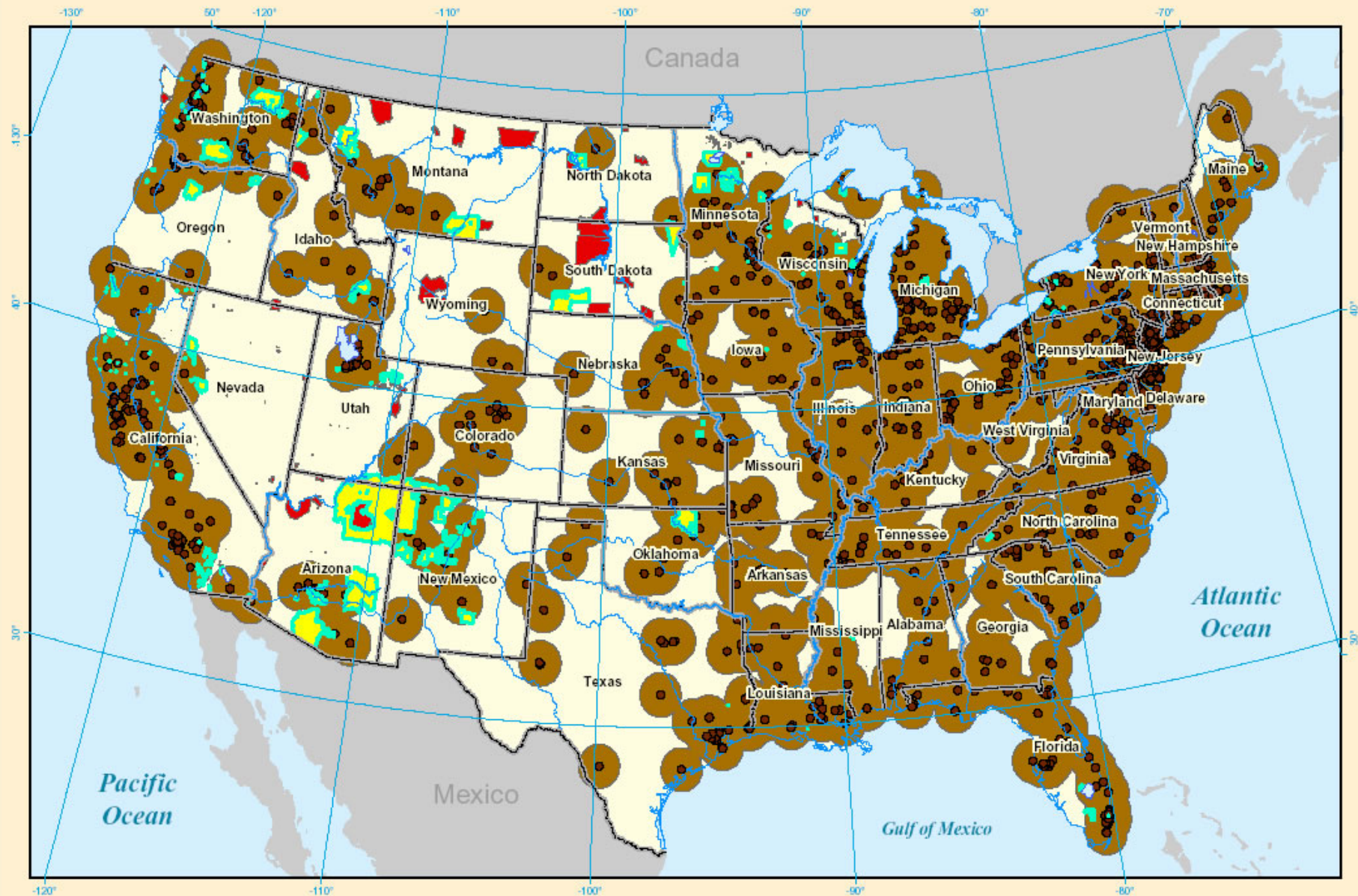
**Indian Lands near
Superfund Sites**



**SBRP Case
Study Areas**



**Kumeyaay Historic
Map by Mike Connolly**



- Reservations near Superfund sites
- Indian Reservations
- Superfund Sites
- 50 miles buffer of Superfund sites

Tribal Lands and Superfund Sites

319 out of 542 Indian Reservations are within 50 miles from Superfund Sites

0 50 100 200 300 Miles

UCSD SBRP Outreach Team
March 2004

Sources: ESRI Data & Maps CD
US Federal Lands database
EPA CERCLIS database



- Tour of Dr. Robert Tukey's lab for tribal science partners
Thursday, July 28, 2005
Dr. Marshall Cheung, Director 29 Palms Tribal science lab



Endocrine disruptor study



Visit to 29 Palms Tribal Science Lab



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Research Translation Core

Dr. Keith Pezzoli, Core Leader

Keith Pezzoli, Ph.D.

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e-mail: kpezzoli@ucsd.edu

web: <http://regionalworkbench.org>

Applying toxicogenomics and biomolecular technologies to environmental monitoring, risk assessment and bioremediation

SBRP Research Translation Core proposal text ([pdf file, 1.8m](#))

SBRP Research Translation Core letters of support ([pdf file, 21.0m](#))



**3 required
components**



SBRP network



SBRP science



**SBRP org
chart**

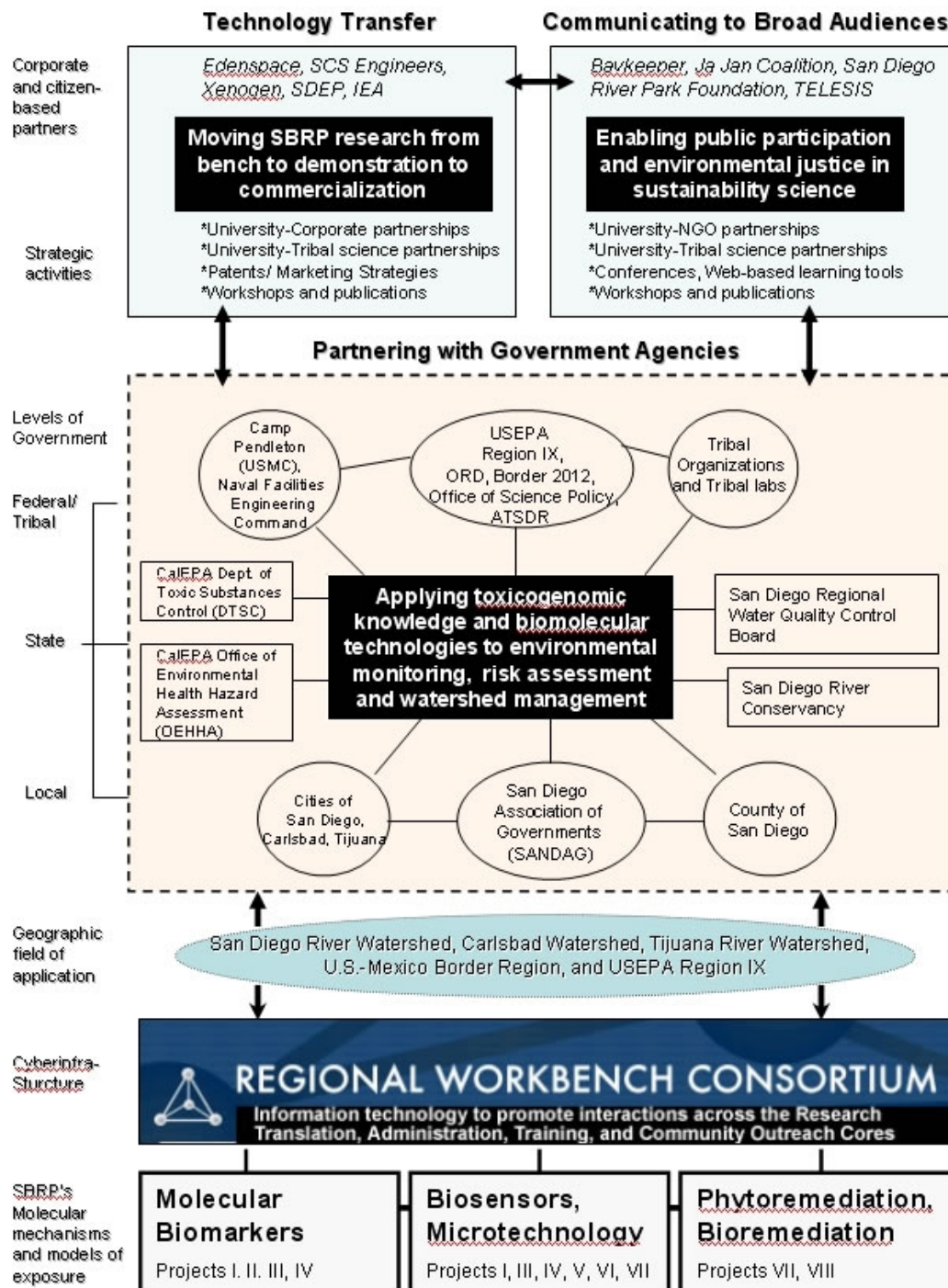


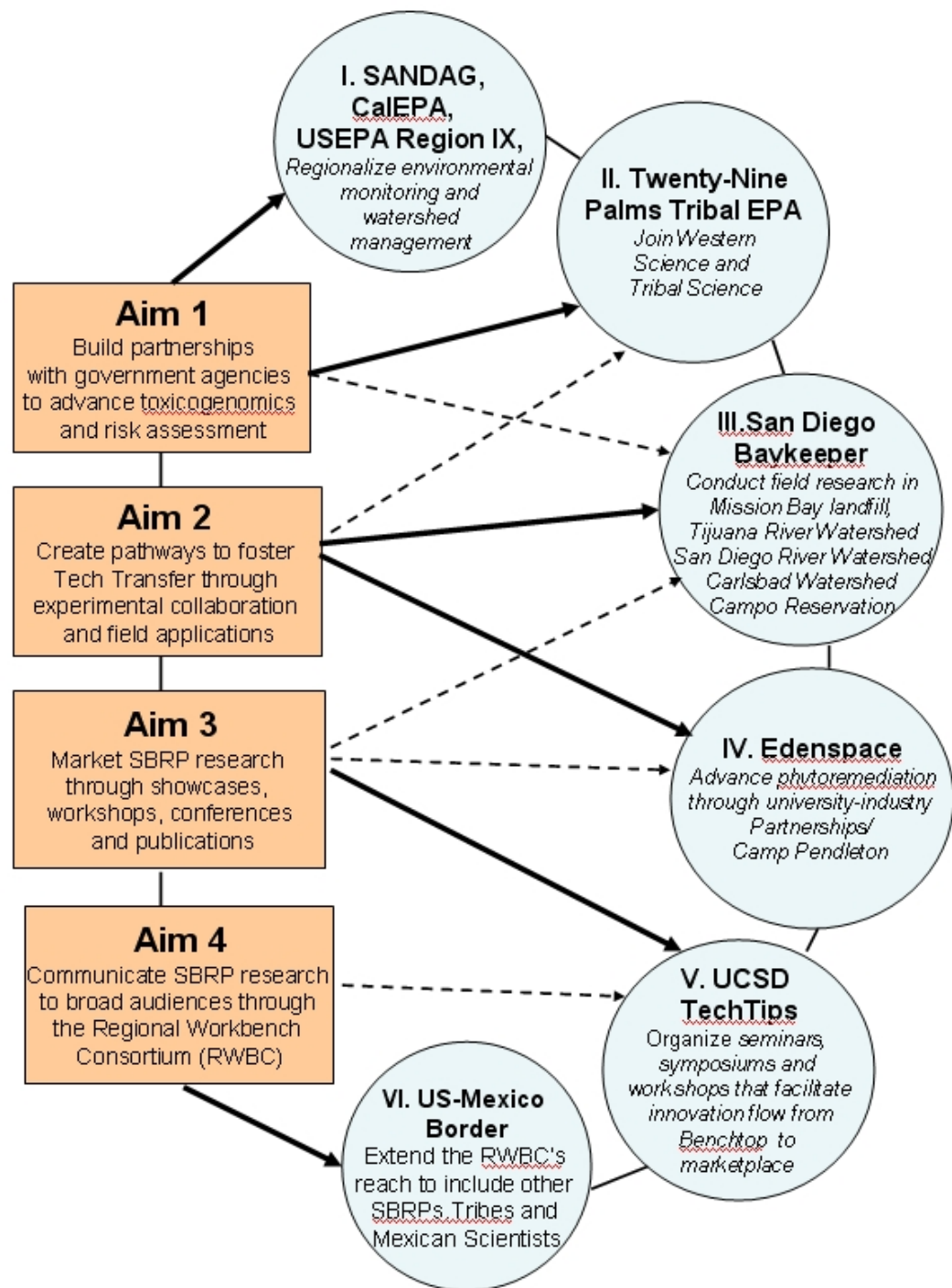
**Research
Translation
Core**



**Aims and
activities**

A selection of images from the proposal, click thumbnail for larger version.





Science, Standards and the TMDL Process In The Urban Watershed Context



GEOGRAPHICAL
INFORMATION SYSTEM





Hiram Sarabia

Citizen Science: Training Session



Water Quality Sampling Methods Training

OK, so you're ready to devise an evaluation plan for your research translation and/or outreach core.

Now what?

1. Set the agenda and participatory incentive structure (what gets on your radar screen? who gets to put it there, how? why?)
2. Operationalize your measures of success (define your metrics, qualitative/quantitative indicators) and methods for ongoing data collection.
3. Listen to learn
4. Interpret the results, archive findings and modify your approach accordingly

1. Set the agenda and participatory incentive structure (what gets on your radar screen? who gets to put it there, how? why?)

Establish a purposeful context and inclusive process that is mutually rewarding for those who you need/expect to participate.

- Use the RFA's conceptual architecture
- Relate your work to EPA/ATSDR
- Identify initial targets in collaboration with your strategic partners including the scientists (to do this well you might first need to go through a “vision and requirements” clarification process).
- Define an incentive structure (figure out how to motivate participation in the process; be entrepreneurial).

US EPA Region IX, Strategic Plan, 2004	<u>Select Goals aligned with the US EPA's 2003-2008 Strategic Plan</u> <ul style="list-style-type: none"> •Improve Water Quality on a Watershed Basis (Aim 1 & 2) •Improve Coastal and Ocean Waters (Aim 1 & 2) •Cleanup and Reuse Contaminated Land (Aim 2) •Reduce Exposure to Toxic Pesticides (Aim 2) •Sustain and Restore US-Mexico Border Ecosystems (Aim 1 & 4) •Build Tribal Capacity (Aim 1 & 4) •Improve Environmental Information Systems (Aim 4) •Engage in joint work with CalEPA and US EPA (including Water Program Issues, Mexican Border, Information Management, and Environmental Indicators) (Aim 1)
ATSDR Goals, accessed on-line, May 2004	<ul style="list-style-type: none"> •Evaluate human health risks from toxic sites and releases and take action in a timely and responsive public health manner (Aim 2) •Ascertain the relationship between exposure to toxic substances and disease (Aim 2) •Develop and provide reliable, understandable information for people in affected communities and Tribes, and stakeholders (Aim 4) •Build and enhance effective partnerships (Aim 1 & 4)

Translational Research

Community Outreach

SBRP Projects	New Biological Models and Technologies	Applications
5 biomedical and 2 non-biomedical projects	<i>Aimed at understanding the impact of Superfund hazardous substances on cellular signaling mechanisms, toxicity, metabolism, endocrine function and overall physiology.</i>	<i>New and improved biological models and technologies for hazardous substance detection, assessment, evaluation, and remediation.</i>
Molecular Biomarkers		
Karin (1) Russell (2) Evans (3) Tukey (4) Tebo (8)	<ul style="list-style-type: none"> Genetically modified mice that are highly sensitive to oxidative stress as well as non-genotoxic and genotoxic hepatocarcinogens. Strains of <i>S. pombe</i> that are sensitive to oxidative stress response. Transgenic mice sensitive to PXR and CAR receptor ligands. New cell lines that can detect xenobiotic receptor activators, Ah-receptor ligands, and arsenic. Biological reagents to detect exposure to complex mixtures of hazardous substances. 	<ul style="list-style-type: none"> Model transgenic and genetically altered organisms (yeast, mice, cell-based systems) useful for risk assessment. New biological methods for detecting/testing toxicants in water/soil/sediment samples. New methods to assess the risk associated with exposure to mixtures of toxicants (e.g., identification of “signature” patterns of particular mixtures). Development of biomarkers for heavy metal exposure and bioavailability (specifically hexavalent Cr, and, possibly Pb and Cu)

Confirm your labs interest by placing a check next to one of these applications. And list below a contact (yourself or someone else) that will serve as liaison between your lab and the RT and CO Cores.

Translational Research

Community Outreach

SBRP Projects	New Biological Models and Technologies	Applications
Biosensors and Microtechnology		
Karin (1) Evans (2) Tukey (3) Taylor (6) Schroeder (7)	<ul style="list-style-type: none"> •Microscale liver tissue modeling technologies. •Bioengineered ('biomimetic') lab-on-a-chip platforms. •New detection methods for monitoring organophosphate inhibited acetylcholine esterase. Transgenic plants 	<ul style="list-style-type: none"> •Reduce animal-animal variability associated with in vivo experiments •Bioengineered chips to explore new technologies for rapid screening of environmental toxicants •Fieldable biosensors to detect exposure to pesticides and heavy metals.
Phytoremediation/Bioremediation		
Schroeder (7, Tebo (8)	<ul style="list-style-type: none"> •Transgenic plant technology for phytoremediation •Engineered organisms (aerobic and anaerobic) for enhanced bioremediation of heavy metals. 	<ul style="list-style-type: none"> •Phytoremediation of heavy metal contaminated soils. •Microbial detoxification (bioremediation by immobilization) of hexavalent metals. •Bioremediation of heavy metals (e.g., Cr, Pb, Cu, Ni, Zn, Cd, As).
Confirm your lab's interest by placing a check next to one of these applications. And list below a contact (yourself or someone else) that will serve as liaison between your lab and the RT and CO Cores.		

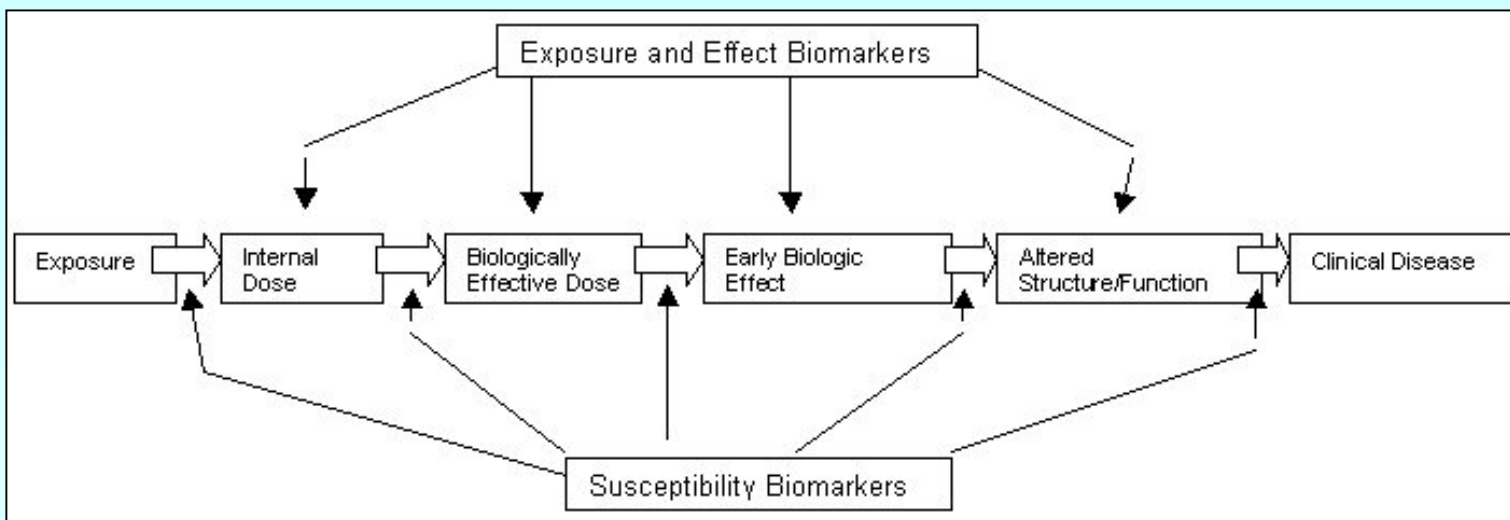


Figure 1 (Adapted from DeCaprio, 1997); cited in **Funding Opportunities Application of Biomarkers to Environmental Health and Risk Assessment** FY 2004 Science to Achieve Results (STAR) Program.

EPA INTEREST in BIOMARKERS

EPA, through the STAR program, is interested in supporting research that provides validation, interpretation and/or application of currently known biomarkers. Of special interest is the use of multiple biomarkers that can fill knowledge gaps across different points of the exposure-dose-effect continuum and/or that can be applied in a clinical setting. Any of the following areas are of interest:

- Mechanistic studies (e.g., using genomics or proteomics) of toxicant response linked to clinical disease. For example, the identification of the functional relevance of proteins where genetic polymorphisms have been found to modify the effect of an environmental exposure on a disease endpoint.
- Studies to validate the utility of biomarkers for use in large population studies (e.g., reliability, predictive value, sensitivity, specificity, affordability, applicability to the general population and susceptible subpopulations).

International Conference: Biomarkers for Toxicology and Molecular Epidemiology, New Tools for 21st-Century Problems

The National Center for Environmental Health/Agency for Toxic Substances and Disease Registry (NCEH/ATSDR) March 15-17, 2004, in Atlanta, Georgia.

Examine ways that cutting-edge biotechnology/molecular tools may be applied to several public health problems through technology transfer. Discuss the latest developments in biosensor and other new analytical technologies **that can be utilized** for rapid, field-usable exposure assessments of environmental chemicals as well as for chemical terror agents.

Evaluate the latest developments in the application of molecular "omics" technologies (genomics, proteomics, and metabolomics). These biomarker classes can be used to **evaluate** early responses to toxic agents such as arsenic in drinking water, which is a global public health problem that affects the United States and many other countries.

Examine these new molecular tools in relation to populations at special risk for toxicity from chemicals (e.g., children). **Attendees will seek to determine ways that these modern tools can be used to provide improved risk assessments for protecting children against environmental chemicals.**

2. Operationalize your measures of success (define your metrics, qualitative/quantitative indicators) and methods for ongoing data collection.
- Quantify your SBRP's output/contributions in the categories listed by Dr. Wilson and Dr. Suk (e.g. publications, awards, patents, service to the EPA).
 - Do your homework and find studies that include indicators or metrics relevant to your aims.
 - Leverage related research and educational efforts on your campus (e.g., ICT and sustainability science, policy and planning, curriculum mandated service learning and field research requirements) to build in an ongoing data collection system.

Biomarker-Based Analysis for Contaminants in Sediments/Soil: Review of Cell-Based Assays and cDNA Arrays (ERDC TN-DOER-C19, December 2000)

Cell-based biomarker assays and cDNA arrays have the potential to be rapid, sensitive, and low-cost tools for sediment/soil toxicity screening.

The P450RGS assay illustrates the potential savings and benefits offered by cell-based assays. Although gas chromatography-mass spectroscopy (GC-MS) is the definitive method for identifying and determining the precise amount of dioxin and dioxin-like compounds present in an environmental sample, the method is expensive (~\$1,200 - \$2,000 per sample). **P450RGS is far less expensive at about \$200 per sample, and its use can result in major cost savings** when used to screen multiple sediment cores to determine which samples should be confirmed by GC-MS.

Inouye, L. S., and McFarland, V.A. (2000). "Biomarker-based analysis for contaminants in sediments/soil: Review of cell-based assays and cDNA arrays," *DOER Technical Notes Collection* (ERDC TN-DOER-C19), U.S. Army Engineer Research and Development Center, Vicksburg, MS. www.wes.army.mil/el/dots/doer
<http://el.erdc.usace.army.mil/dots/doer/pdf/doerc19.pdf>

Senior Sequence

Urban Studies & Planning

University of California, San Diego

News & Updates



NEWS AND KNOWLEDGE NETWORKING

[Senior Sequence News and Announcements blog](#)

[Senior Sequence Areas of Concentration Blog](#)



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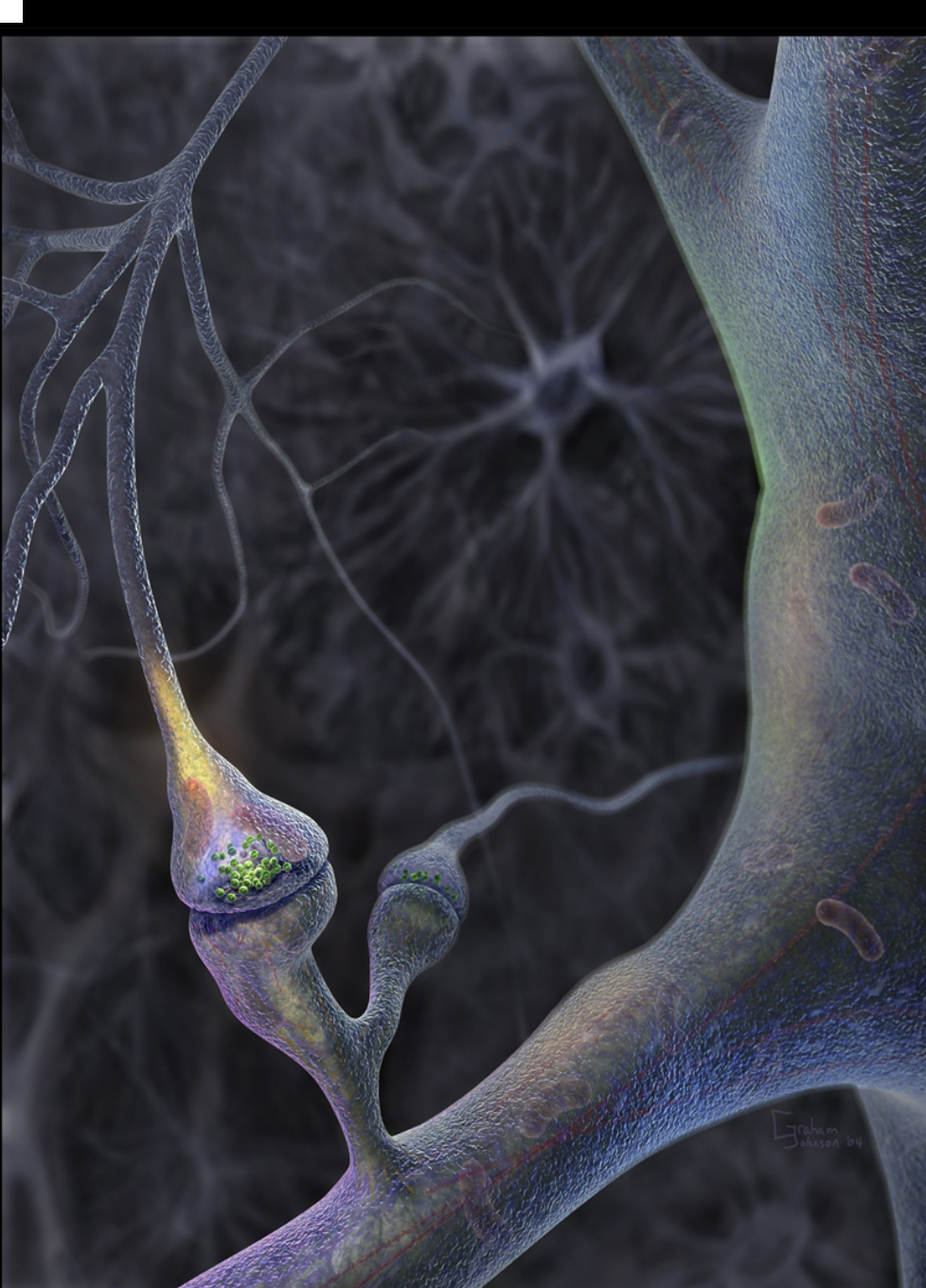
3. Listen to learn (EPA)

Exploit the real power of evaluation (go beyond a mere number crunching exercise to justify funding levels).

Use the process to foster mutual learning and build social capital (enriched partnerships, knowledge networks and communication systems).

4. Interpret the results, archive findings and modify your approach accordingly

- Enrich your interpretative framework by incorporating insights from key disciplines/literature/methods in social science, information science and the humanities (organizational culture, social network analysis, new institutionalism, knowledge management).
- Create an archive of lessons learned, best practices that can be shared.



Suggested Next Steps:

- Enhance knowledge networking capacity for SBRP programs to **“federate distributed intelligence”** in the effort to link environmental health sciences to society for the common good--for example:
 - 1. Create a national database of best practices, literature, success/failure stories in evaluation methods/practice specific to SBRP research translation and community outreach
 - 2. Conduct a professionally mediated workshop (on evaluation training and standards) for all PIs and key staff of the RTCs and COCs

Contact info

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SBRP, Research Translation/
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